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Abstract: **PURPOSE** The aim of this systematic review and meta-analysis was to evaluate the microbial colonization at the implant-abutment interfaces (IAI) on bone-level implants and to identify possible association with peri-implant conditions. **STUDY SELECTION** The focus question aimed to answer whether two-piece osseointegrated implants, in function for at least 1 year, in human, relate to higher bacterial count and the onset of periimplantitis, compared to healthy peri-implant conditions. Search strategy encompassed the on-line (MedLine, Google scholar, Cochrane library) literature from 1990 up to March 2015 published in English using combinations of MeSH (Medical Subject Headings) and search terms. Quality assessment of selected full-text articles was performed according to the ARRIVE and CONSORT statement guidelines. For data analysis, the total bacterial count of *Porphyromonas gingivalis*, *Tannerella forsythia*, *Treponema denticola*, *Prevotella intermedia*, and *Fusobacterium nucleatum* was calculated and compared to IAI with or without peri-implant pathology. **RESULTS** A total of 14 articles, reporting data from 1126 implants, fulfilled the inclusion criteria and subjected to quality assessment. The selected studies revealed contamination of the IAI, in patients who received two-piece implant systems. Meta-analysis indicated significant difference in total bacterial count between implants affected by periimplantitis versus healthy peri-implant tissues (0.387 ± 0.055 ; 95% CI 0.279-0.496). Less bacterial counts were identified in the healthy IAI for all the investigated gram-negative bacteria except for *T. forsythia*. **CONCLUSIONS** Significantly higher bacterial counts were found for periodontal pathogenic bacteria within the IAI of implants in patients with periimplantitis compared to those implants surrounded by healthy peri-implant tissues.

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Microbial Colonization at the Implant-Abutment Interface and Influence on the Onset of Peri-Implantitis: A Systematic Review and Meta-Analysis

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Authors' contribution

Tallarico M.: substantially contributed to conception and design; contributed to acquisition, analysis, and interpretation of data; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

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Abstract

Purpose: The aim of this systematic review and meta-analysis was to evaluate the microbial colonization at the implant-abutment interfaces (IAI) on bone-level implants and to identify possible association with peri-implant conditions.

Materials and Methods: The focus question aimed to answer whether two-piece osseointegrated implants in function for at least 1 year in human relate to higher bacterial count and the onset of peri-implantitis, compared to healthy peri-implant conditions. Search strategy encompassed the on-line (MedLine, Google scholar, Cochrane library) literature from 1990 up to March 2015 published in English using combinations of MeSH (Medical Subject Headings) and search terms. Quality assessment of selected full-text articles was performed according to the ARRIVE and CONSORT statement guidelines. For data analysis, the total bacterial count of *Porphyromonas gingivalis*, *Tannerella forsythia*, *Treponema denticola*, *Prevotella intermedia*, and *Fusobacterium nucleatum* was calculated and compared to IAI with or without peri-implant pathology.

Results: A total of 14 articles, reporting data from 1126 implants, fulfilled the inclusion criteria and subjected to quality assessment. The selected studies revealed contamination of the IAI, in patients who received two-piece implant systems. Meta-analysis indicated significant difference in total bacterial count between implants affected by peri-implantitis versus healthy peri-implant tissues (0.387 ± 0.055 ; 95% CI 0.279-0.496). Less bacterial counts were identified in the healthy IAI for all the investigated gram-negative bacteria except for *Tannerella forsythia*.

Conclusions: Significantly higher bacterial counts were found for periodontal pathogenic bacteria within the IAI of implants in patients with peri-implantitis compared to those implants surrounded by healthy peri-implant tissues.

Introduction

Microgaps at the implant-abutment interfaces (IAI) are typical for two-piece osseointegrated dental implant systems and seem to play a significant role in bacterial colonization at the peri-implant sulcus.¹ This, in turn, may yield to peri-implant inflammatory reactions and subsequently loss of supporting bone.²⁻⁷ Bacterial leakage at the IAI along with the abutment screw assemblies that act as bacterial reservoir may result in an area of the fixture/abutment interface and trigger a host response with inflamed soft tissues and possible marginal peri-implant bone loss.⁸⁻¹²

Numerous attempts have been made to reduce the inner bacterial colonization at the IAI, among which the application of 0.2% chlorhexidine solution at two stage surgeries is considered a more common practice. Yet, controversial opinions exist on the effectiveness of chlorhexidine solution in preventing microbial colonization at the IAI.^{13,14} Bacterial endotoxins typically penetrate the IAI especially with Morse-taper connection but 0.2% chlorhexidine solution could not significantly eliminate the penetration. Alternative cleaning method, such as plasma of argon, was claimed to be a favorable method to reduce the inner bacterial colonization at the IAI, maintaining hard tissue levels.⁷

The objectives of this systematic review and meta-analysis were to evaluate the microbiological colonization at the implant-abutment interface on bone level implants and investigate whether it relates to the onset of peri-implantitis.

Materials and Methods

This systematic review conformed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines (<http://www.prisma-statement.org>).¹⁵ The protocol of this systematic review has been published in the international prospective register of systematic reviews (PROSPERO, <http://www.crd.york.ac.uk/PROSPERO/>) with registration number CRD42016037481. The focused question of the review was to identify

whether there is a relationship between the presence of higher bacterial count and the onset of peri-implantitis, compared to healthy peri-implant conditions in patients with two-piece osseointegrated implants after at least 1 year of function. Peri-implantitis was defined by the presence of peri-implant probing depth ≥ 5 mm associated with bleeding on probing and/or suppuration, and radiographic images of bone loss ≥ 3 mm, compared to initial radiographs at delivery of the prosthetic restoration.^{16,17}

Information Sources

Articles published only in English were searched that reported on microbial colonization at the IAI and its relationship with the onset of peri-implantitis, published from 1990 until March 2015 PubMed database of the US National Library of Medicine (<http://www.ncbi.nlm.nih.gov/pubmed/>), Google scholar (<http://www.google.com>) and the Cochrane Library (<http://www.cochranelibrary.com/>). Furthermore, the references of the included articles were checked manually in order to find additional articles.

Search Strategy

Initially, PICOS question (Population (P), Intervention (I), Comparison (C), Outcomes and Study Design (O), Study type (S)) defined the search strategy, where P=Two-piece osseointegrated implants with a diagnosis of peri-implantitis after at least 1 year of function; I=Microbial colonization at the IAI; C=Healthy peri-implant conditions; O=Survival rate; S=Randomized controlled clinical trials (RCT) and clinical follow-up studies.¹⁸

The electronic databases were searched using a combinations of MeSH (Medical Subject Headings) terms, search terms and their combinations: "dental implants" [MeSH] AND "bacterial contamination" OR "presence of bacterium" OR "dental leakage/microbiology" [MeSH] OR "microleakage" OR "microbiological findings" OR "microbiological colonization" OR "microbiota" OR "peri-implant microflora" AND "peri-implantitis" [MeSH] OR "peri-implant pathology" OR "peri-implant disease" AND "Dental Abutments*/microbiology" [MeSH] "connection, implant-abutment" OR "dental Implant-abutment design" [MeSH] OR

“implant-abutment junction” OR “implant-abutment microgap” OR “inner space of dental implants” OR “inner part of dental implants”.

Study Selection and Eligibility Criteria

All titles and abstracts of the selected studies were first assessed for the following inclusion criteria: 1) Articles written in English; 2) Studies with a clinical examination of the patients; 3) Studies assessing the counts of different bacterial species (bacterial count, BC) at the IAI level in patients who received two-stage bone level implant systems, independently from the configuration of the connection; 4) Randomized controlled clinical trials (RCTs), prospective cohort studies or cross-sectional studies reporting on implants in function for at least 1 year.

After evaluating the full text of the articles according to the previously defined exclusion criteria, articles with the following features, without restriction in languages, were not considered eligible: a) Letters, narrative or historical reviews; 2) Animal and in vitro studies; 3) Reports on locally or systemically compromised sites and/or conditions (i.e. major bone defect before implantation, bone pathologies, head and neck radiotherapy, treatment with bisphosphonates); 4) Reports on patients who received mechanical debridement in the previous 3 months or antibiotics in the last 6 months before analysis.

Data Collection Process

Two calibrated reviewers (M.C. and L.C.) screened and collected the data from selected papers onto structured tables. Cohen`s Kappa values between examiners was calculated at both the first and the second stage of the research. Discrepancies were resolved by consensus and a third examiner (M.T.) was consulted.

Articles without abstracts but with titles related to the objectives of this review were selected and their full text were screened for eligibility. Reference lists of the selected articles were further screened for possible additional papers. Additionally, hand searches of the bibliographies of selected systematic reviews were conducted limited to the

following journals: Clinical Implant Dentistry and Related Research; Clinical Oral Implants Research; International Journal of Oral and Maxillofacial Implants; Journal of Clinical Periodontology; Journal of Periodontology.

Assessment of quality, heterogeneity and Risk of Bias of Individual Studies

The same reviewers assessed the risk of bias in the included sample according to the guidelines provided by the CONSORT statement for the evaluation of randomized controlled trials (<http://www.consort-statement.org>), the STROBE statement for observational studies (<http://www.strobe-statement.org>), as well as the modified items from the Cochrane Collaboration Tool for assessing risk of bias (Table 1).^{19,20}

Considering the adequacy in the respective studies, the items were graded and the percentage of positively graded items was calculated.¹⁹ Quality assessment was performed in two different phases, namely phase I where quality assessment was based on the published full-text articles performed independently by both reviewers and in phase II where disagreements were resolved upon discussion. After collecting the scores at phase II of quality assessment, an overall estimation of plausible risk of bias (low, moderate or high) was completed for each selected study. While a low risk of bias was estimated when all the criteria were met, a moderate risk was considered when one or more criteria were partly met and a high risk of bias was estimated when one or more criteria were not meet (Cochrane Handbook for Systematic Reviews of Interventions, version 5.1.0. [http:// www.cochrane.org/resources/handbook](http://www.cochrane.org/resources/handbook)).

Measures and Analysis of Results

Descriptive statistics, meta-regression and meta-analysis were performed, based on the comparable studies reporting the same outcome measures. The microbiota present at the IAI of implants in function for at least 1 year was considered for data analysis. BCs of gram-negative bacteria associated with chronic periodontitis (*Porphyromonas gingivalis*,

Tannerella forsythia, Treponema denticola, Prevotella intermedia, and Fusobacterium nucleatum) were extracted and defined as primary outcome variable.²¹ The meta-regression considered microbiota that are regularly detected at peri-implantitis sites and are found to increase the risk for peri-implant bone loss and disease progression.^{10,21-23} Mean differences were combined using random-effects models. Heterogeneity between studies, subgroup analyses, meta-analysis, and forest plots were calculated using a software program (Comprehensive Meta-Analysis V3; Biostat, Englewood, NJ, USA).

Results

Study Selection

A total of 523 potentially relevant titles and abstracts were found after the electronic and manual search. During the first stage of selection, 309 articles were excluded based on the titles and abstracts ($k=0.72$). During the second phase, complete full-text articles of the remaining 212 publications were evaluated and 198 articles were excluded since they did not fulfill the inclusion criteria ($k=0.98$). Finally, a total of 14 articles, reporting data from 1126 implants, were selected that fulfilled inclusion criteria and quality assessment required for this systematic review (Fig. 1).

Study Characteristics

The 14 selected articles were published between 1993 and March of 2015, two of which were RCTs,^{24,25} two prospective cohort studies,^{26,27} and ten cross-sectional studies.^{5,7,28-35} Only one prospective clinical study (27) was written following the STROBE statement for observational studies (<http://www.strobe-statement.org>). Hence, a direct comparison between the selected articles was not possible.

Risk of Bias within Studies

One publication was associated with a low risk of bias,²⁷ seven with moderate risk of bias,^{5,7,32-35} and six with high risk of bias.^{25,26,28-31}

The included articles received minimum grading when evaluating submission to ethical committees (6/14), presence of blinded evaluators (2/14), standardization of the procedures (1/14) and presence of eligible criteria (9/14) (Table 1).

Measures and Meta-regression Analysis

Bacterial leakage at the IAI: All selected studies reported contamination of the IAI and the abutment surface in patients receiving the assembly of a two-stage implant system. Quantitative real-time polymerase chain reaction (PCR) was carried out for BC in 7 of the 14 studies,^{5,7,24,27,33,34} where the following pathogens were analyzed: *Aggregatibacter actinomycetemcomitans*, *Porphyromonas gingivalis*, *Tannerella forsythia*, *Treponema denticola*, *Prevotella intermedia*, *Parvimonas micra*, *Fusobacterium nucleatum*, *Campylobacter rectus*, *Eikenella corrodens*, *Candida albicans*, *Enterococcus faecalis*, and *Porphyromonas aeruginosa*. While in one study the checkerboard DNA-DNA hybridization technique was used,³² in other six studies different techniques including a scanning electron microscopy was used in order to screen the colony morphology.^{25,26,28-31} In one study,²⁷ progressive colonization by periodontal pathogenic bacteria was described in the internal portions of two-piece implants. In another study,³² intra-coronal components of screw-retained fixed restorations were heavily contaminated in all the specimens. Contamination of abutment screws most likely occurred from the peri-implant sulcus through the IAI and abutment-prosthesis interface. Likewise, significant differences in antibiotic-resistant nosocomial bacteria (*E. faecalis* and *P. aeruginosa*) were observed at the internal and external implant components between healthy peri-implant sulci and implants compromised with peri-implantitis.³⁵ Regarding the absence/presence of the bacteria analyzed, no relevant differences were found between the analysis at the peri-implant sulcus and the connections inside the abutments surfaces.⁵ The microbial composition at the neighbouring teeth resembled those found in the the peri-implant

sulcus with a high frequency for *P. gingivalis*, *T. forsythia*, *P. intermedia*, *P. micra* and *E. corrodens*.⁵

Two comparative studies between healthy peri-implant conditions versus implants affected by peri-implantitis,^{5,7} reported bacterial contamination in both groups. Orange complex species (*P. intermedia*, *P. micra*, *F. nucleatum*)^{36,37} were the most prevalent in all sites analyzed for both groups. Inside of the implant connection, the prevalence of the analyzed species was more predominant in the peri-implantitis group and varied from 1.1% *A. actinomycetemcomitans* to 98.9% *F. nucleatum*. Species with $\geq 50\%$ of prevalence were: *P. gingivalis*, *T. denticola*, *P. intermedia*, *F. nucleatum*, *C. rectus*, *E. corrodens*, *T. forsythia* and *P. micra*.⁵

Bacterial leakage at the IAI in relation to abutment connection design: The selected sample showed greater heterogeneity regarding the type of the IAI. Four studies reported on external hexagon connections,^{24,28,30,32} and two studies either on internal hexagons²⁷ or morse taper²⁹ connections. Four studies used different IAI designs,^{5,25,31,34} while the type of IAI was not reported in the other 4 manuscripts.^{7,26,33,35}

The evaluation of four different IAIs implied that all the analyzed connections presented contamination after 5 years of functional loading.³⁴ It also appeared that the connection design might have influenced the BC levels qualitatively and quantitatively, especially inside the implant connections, showing better results for the conical connection. Similarly, different types of abutments showed significant variation on the mean microgap size within the first 5 hours of loading.²⁵ However, no significant influence of micro-leakage was found at 24 hours, 48 hours, and 14 days on BC levels. Yet, the use of standard abutments significantly decreased the microgap size compared to customized ones. The study concluded that the microleakage in the connection area was comparable for all of the analyzed abutments.

Meta-regression and Analysis of Subgroups

Five studies, including a total of 622 implants (n=223 with peri-implantitis; n=399 with healthy peri-implant conditions) in function for at least 1 year, were included in the meta-analysis (34; 27; 33; 5; 7). BC of gram negative anaerobic showed relevance to chronic periodontitis and founded to increase the risk for peri-implant bone loss and disease progression due to the presence of periodontal pathogens (*P. gingivalis*, *T. forsythia*, *T. denticola*, *P. intermedia*, *F. nucleatum*) (Table 2).^{10,21-23} Meta-analysis considered the bacteria that were evaluated in all selected studies. Two out of five studies compared the BC in healthy peri-implant conditions versus implants affected by peri-implantitis (5; 7). Meta analysis revealed higher mean values for the BC of all the gram-negative bacteria analyzed, except for *T. forsythia* in implants with peri-implantitis (Fig. 2). Overall, the mean differences in BC were statistically significant between the two analyzed groups, with higher values in implants with peri-implantitis (difference: 0.387 ± 0.055 ; 95% CI 0.279-0.496, $p=0.000$).

Discussion

This systematic review evaluated the microbial colonization at the IAI on bone level implants and related it to the possible onset of peri-implantitis. Except for *T. forsythia*, significantly higher BC was identified at implants affected by peri-implantitis compared to healthy peri-implant sulci for all gram-negative plausible periodontal pathogens. For the *T. forsythia*, only a trend towards higher BC was detected.

The included studies assessed the microbiota at the level of IAI in patients who received two-stage bone level implant systems with various implant-abutment connection designs. Nevertheless, two studies reported the contamination of the IAI independent from the connection design.^{25,34} Furthermore, no distinction could be made between screw- and cemented-retained restorations assuming that the crown-abutment junction is located more coronal, assuming that the gaps is filled with cement.

Presence of bacterial contamination at the IAI placed at the alveolar bone level was demonstrated to be associated with significant inflammatory cell infiltration and bone loss.²⁷ Increased accumulation of acute inflammatory cells adjacent the IAI suggests the persistence of chemotactic stimuli from this region sustaining continuous recruitment of neutrophilic granulocytes.^{38,39} Additionally, the presence of an inflammatory infiltration of peri-implant tissue at the fixture-abutment interfaces was also led by microbial internal contamination.^{24,26}

A number of studies reported that microbial contamination could occur at the level of IAI both in implants with healthy and diseased tissue conditions.^{5,7,41} Despite the fact that there were no clinical signs of peri-implantitis, the presence of the bacterial species associated with this condition were clearly elevated. When clinical and microbiological characteristics in subjects and implants with healthy tissue conditions or peri-implantitis were evaluated and data from healthy and diseased implant sites were compared within the same subject (534 patients; 1507 dental implants), clear trends were observed.⁵ Microbial analysis obtained from three locations (peri-implant sulcus (PIS), inner parts of the implant connections (PI), gingival sulcus of the neighbouring teeth) along with clinical parameters (bleeding on probing, probing pocket depth, plaque index), presence of peri-implantitis was evident in 10.3% of the patients and in 7.3% of the implants. The microbial analysis within the 53 patients affected by peri-implantitis revealed no relevant differences between the analysis at the PIS and PI.⁵

Microgap at the IAI may also yield to mechanical and biological complications including abutment screw fractures and peri-implant diseases.²⁵ Microgap size and microbial leakage at different times at the IAI of 4 different abutments to Straumann implants denoted significant effect on the mean microgap size ($p < 0.001$) and on the mean number of bacterial colonies (CFU/mL) leaking from the IAI within the first 5 hours of the

experiment ($p=0.012$).²⁵ However, the micro-leakage at 24 hours, 48 hours, and 14 days was no longer influenced significantly ($p=0.145$).

Clinical and microbial differences between healthy peri-implant conditions and peri-implantitis revealed that the microbial prevalence was higher in the peri-implantitis group at three locations and the differences in prevalence between different types of bacteria were more marked inside the connection than in the PIS (57 patients; 122 implants).⁵ When opportunistic pathogens (*E. faecalis*, *P. aeruginosa*) were identified in the presence of peri-implant disease at the level of PIS of each implant, gingival sulcus of the adjacent teeth and the connection and abutment at the inner portion of each implant, significant differences on the presence and amount of nosocomial bacteria were detected around diseased implants.³⁵ Not only microbial leakage through the gap between the supra-structure and the abutment,^{27,29} but also implant designs and materials may affect the potential risk of harboring oral microorganisms.^{41,42} Typically morse taper connections seems collect less bacteria as opposed to three-channel connection or conical connection.⁴³ Similarly, morse taper connections presented favorable results in this respect compared to the trilobe cemented connection.⁴⁴ On the other hand, bacterial microbiota present inside the implant connection and in the PIS fluid of implants with healthy peri-implant conditions with four different implant systems after at least 5 years of functional loading, demonstrated microbiological contamination in all types of connections regardless of the site (peri-implant sulcus, inner portion of the connection, abutment surface and gingival sulcus of neighbouring teeth).³⁴

Implications for Clinical Practice

This meta-analysis indicated that bacteria could easily be colonized at the implant-abutment interface and may consequently cause outbreak of peri-implantitis. It is evident from a clinical point of view that inner portions of IAI should always be considered contaminated. Clinicians should note that there exists gap in the clinical evidence for justification

of cleaning IAI at regular intervals to improve soft and hard tissue healing. Yet, current evidence may suggest removal of the crown/abutment complex and the disinfection/sterilization of the connection units both at the implant and abutment aspects at certain intervals as an adjunct to maintenance regimens of dental implants.

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Legends to figures and tables:

Figures:

Figure 1. Flow diagram with the information through the phases of study selection based on PRISMA .

Figure 2. Forest plot of pooled studies and meta-analysis of the differences in bacterial count for each investigated bacteria present at the implant-abutment interface (IAI) of healthy implants and implants affected by peri-implantitis for at least 1 year.

Tables:

Table 1. Reporting quality of all selected full-text articles (Graziani et al., 2012; Pjetursson et al., 2012).

Table 2. Included studies and reported counts of gram-negative anaerobic bacteria (*P. gingivalis*, *T. forsythia*, *T. denticola*, *P. intermedia*, *F. nucleatum*) associated with chronic periodontitis present at the implant-abutment interface (IAI) of healthy implants and implants affected by peri-implantitis for at least 1 year.

Table 1. Reporting quality of all selected full-text articles (Graziani et al., 2012; Pjetursson et al., 2012).

1 st Author & Year	Study Design	Ethical committee	Criteria for Eligibility	Standardization	Blinded Assessment	Connection	Participants	Number of Implants	TBC methods	Risk of Bias
Cosyn et al. 2009	Cross-sectional	Yes	-	Yes	-	External hex	8	58	DNA-DNA hybridization tech-	Moderate
Jervøe-Storm et al. 2009	Prospective	Yes	Yes	Yes	Yes	Internal hex	66	26	Quantitative real-time PCR	Low
Canullo et al. 2015d	Cross-sectional	Yes	Yes	Not clear	-	Not reported	38	52	Quantitative real-time PCR	Moderate
Persson et al. 1996	Cross-sectional	-	-	-	-	External hex	10	28	Colony morphology on the blood agar plates	High
Canullo et al. 2014	Cross-sectional	-	Yes	Not clear	-	Different types (4)	40	60	Quantitative real-time PCR	Moderate
Canullo et al. 2015a	Cross-sectional	Yes	Yes	Not clear	-	Different types (3)	53	231	Quantitative real-time PCR	Moderate
Paolantonio et al. 2008	Randomized controlled trial	Yes	Yes	Not clear	-	External hex	30	30	DNA Extraction and Polymerase Chain Reaction (PCR) Amplifications	Moderate
Peñarrocha et al. 2014	Cross-sectional	-	Yes	Not clear	-	Not reported	20	43	Quantitative real-time PCR	Moderate
Rimondini et al. 2001	Prospective	-	-	-	-	Not reported	17	17	Scanning electron microscopy and energy dispersive x-ray spectroscopy (EDS) analysis.	High
Canullo et al. 2015c	Cross-sectional	Yes	Yes	Not clear	-	Not reported	110	225	Quantitative real-time PCR	Moderate
Scarano et al. 2005	Cross-sectional	-	-	-	-	Different type (5)	-	272	Observed in normal reflecting light under a Laborlux-S light microscope	High
Rismanchian et al. 2012	Randomized controlled trial	-	-	-	-	Different types (4)	-	36	TSB culture and SEM	High

Quirynen et al. 1993	Cross-sectional	-	Yes	-	-	External hex	9	18	Different phase contrast microscopy (DPCM)	High
Keller et al. 1998	Cross-sectional	-	Yes	-	-	Internal morse taper	30	30	Darkfield microscope	High

Table 2. Included studies and reported counts of gram-negative anaerobic bacteria (*P. gingivalis*, *T. forsythia*, *T. denticola*, *P. intermedia*, *F. nucleatum*) associated with chronic periodontitis present at the implant-abutment interface (IAI) of healthy implants and implants affected by peri-implantitis for at least 1 year.

1st Author & Year	Bacteria	Mean	Standard deviation	Variance	Lower limit	Upper limit	Z-Value	P-Value
Peri-implantitis: 1) Canullo et al., 2015a 2) Canullo et al., 2015c	<i>P. gingivalis</i>	3.377	0.186	0.035	3.012	3.747	18.154	0.000
	<i>T. forsythia</i>	2.388	0.158	0.025	2.079	2.697	15.124	0.000
	<i>T. denticola</i>	2.641	0.185	0.034	2.279	3.003	14.281	0.000
	<i>P. intermedia</i>	4.410	0.192	0.037	4.034	4.786	22.966	0.000
	<i>F. nucleatum</i>	5.60	0.120	0.014	5.364	5.836	46.507	0.000
Total		3.516	0.088	0.008	3.344	3.689	40.049	0.000
Healthy implants: 1) Canullo et al., 2015a 2) Canullo et al., 2015c 3) Penarrocha-Oltra et al., 2014 4) Jervøe-Storm et al., 2014 5) Canullo et al., 2014	<i>P. gingivalis</i>	2.475	0.157	0.024	2.169	2.782	15.815	0.000
	<i>T. forsythia</i>	2.783	0.135	0.018	2.519	2.047	20.648	0.000
	<i>T. denticola</i>	1,685	0.148	0.022	1.395	1.975	11.398	0.000
	<i>P. intermedia</i>	3.543	0.167	0.028	3.217	3.870	21.279	0.000
	<i>F. nucleatum</i>	5.193	0.106	0.011	4.986	5.401	48.977	0.000
Total		2.971	0.075	0.006	2.824	3.118	39.555	0.000

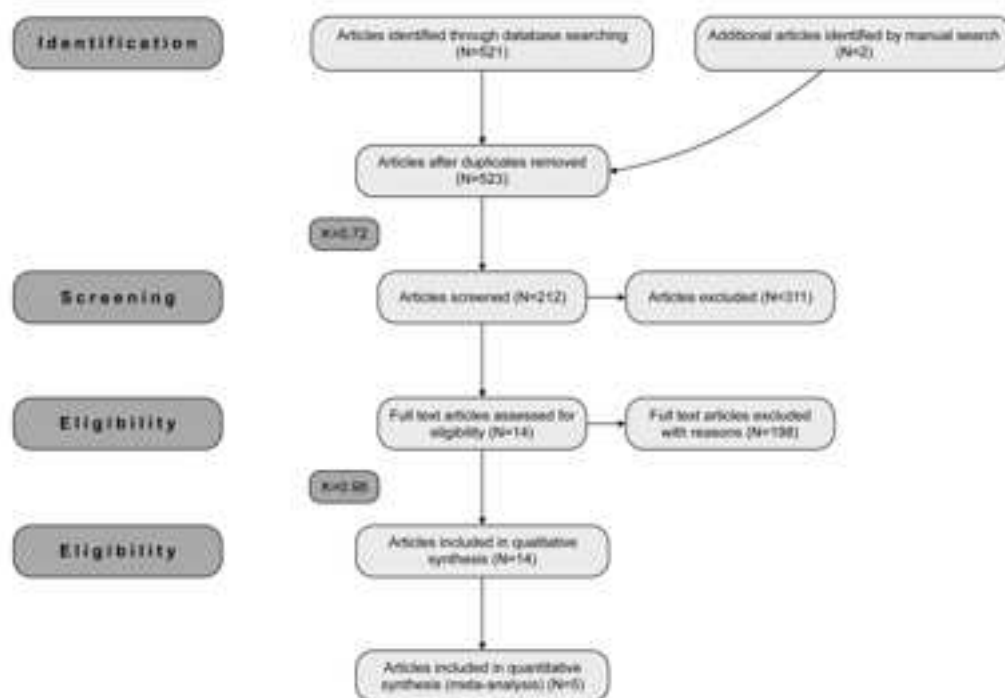


Figure 1

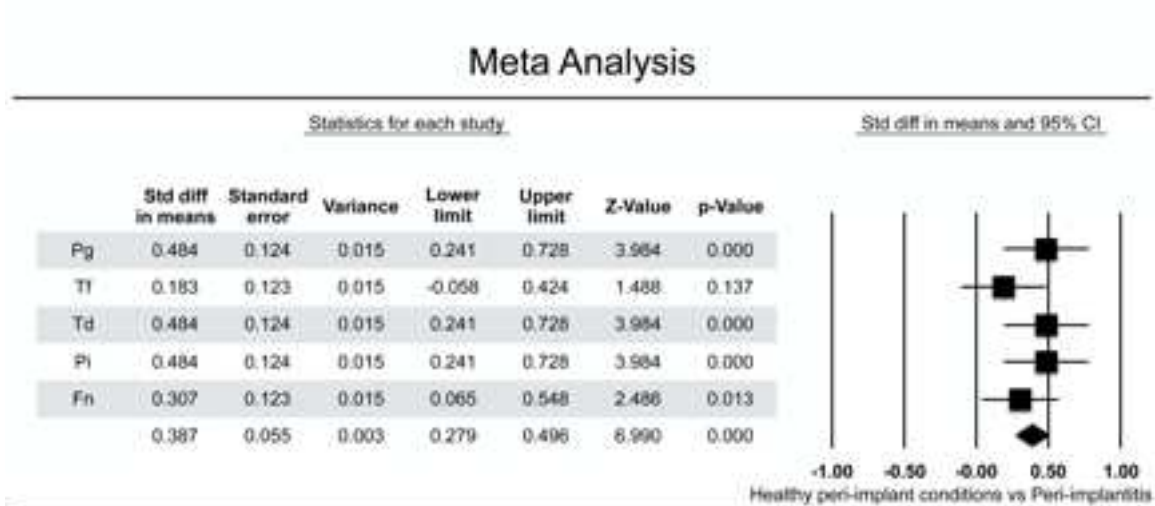


Figure 2